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TRANSMITTAL FORM (to be used for all correspondence after initial filing)	Application Number	10/625124
	Filing Date	07/23/2003
	First Named Inventor	Dobrowolski, Steven et al.
	Art Unit	1645
	Examiner Name	
Total Number of Pages in This Submission	Attorney Docket Number	2263

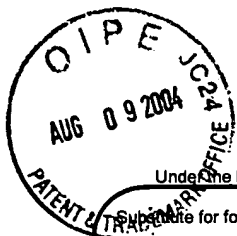
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INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Use as many sheets as necessary)		Complete if Known	
		Application Number	10/625,124
		Filing Date	07/23/2003
		First Named Inventor	Dobrowolski, Steven
		Art Unit	1645
Examiner Name			
Sheet 1	of 1	Attorney Docket Number	2263

NON PATENT LITERATURE DOCUMENTS			
Examiner Initials*	Cite No. ¹	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T ²
		POMPONIO, RJ et al., Profound biotinidase deficiency caused by a point mutation that creates..., Human Molecular Genetics (1997), 6(5), 739-745.	
		COLE, H et al., Human Serum Biotinidase: cDNA Cloning, Sequence, and Characterization, J. Biol. Chem. (1994), 269(9), 6566-6570.	
		HEARD, Gregory et al., A Screening Method for Biotinidase Deficiency in Newborns, Clin. Chem. (1984), 30/1, 125-127.	

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